

**Amendments to the Specification:**

Please amend the specification by replacing the paragraph sections under the heading "Related Applications" with the following new paragraph sections:

**At page 2, lines 34-37 to page 3, lines 1-8:**

R<sup>3</sup> is in the 2-, 3- or 4-position and is:

carboxy; (C<sub>1-6</sub>)alkoxycarbonyl; aminocarbonyl wherein the amino group is optionally substituted by hydroxy, (C<sub>1-6</sub>)alkyl, hydroxy(C<sub>1-6</sub>)alkyl, aminocarbonyl(C<sub>1-6</sub>)alkyl, (C<sub>2-6</sub>)alkenyl, (C<sub>1-6</sub>)alkylsulphonyl, trifluoromethylsulphonyl, (C<sub>1-6</sub>)alkenylsulphonyl, (C<sub>1-6</sub>)alkoxycarbonyl, (C<sub>1-6</sub>)alkylcarbonyl, (C<sub>2-6</sub>)alkenyloxycarbonyl or (C<sub>2-6</sub>)alkenylcarbonyl and optionally further substituted by (C<sub>1-6</sub>)alkyl, hydroxy(C<sub>1-6</sub>)alkyl, aminocarbonyl(C<sub>1-6</sub>)alkyl or (C<sub>2-6</sub>)alkenyl; cyano; tetrazolyl; 2-oxo-oxazolidinyl optionally substituted by R<sup>10</sup>; 3-hydroxy-3-cyclobutene-1,2-dione-4-yl; 2,4-thiazolidinedione-5-yl; tetrazol-5-ylaminocarbonyl; 1,2,4-triazol-5-yl optionally substituted by R<sup>10</sup>; or 5-oxo-1,2,4-oxadiazol-3-yl; or

(C<sub>1-4</sub>)alkyl **optionally substituted** or ethenyl substituted with any of the substituents listed above for R<sup>3</sup> and up to 3 groups R<sup>12</sup> independently selected from:

**At page 4, lines 11-19:**

A is NR<sup>11</sup> or CR<sup>6</sup>R<sup>7</sup> and B is NR<sup>11</sup>, O, SO<sub>2</sub> or CR<sup>8</sup>R<sup>9</sup> and wherein:

each of R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is independently selected from: hydrogen; (C<sub>1-6</sub>)alkylthio; halo; trifluoromethyl; azido; (C<sub>1-6</sub>)alkyl; (C<sub>2-6</sub>)alkenyl; (C<sub>1-6</sub>)alkoxycarbonyl; (C<sub>1-6</sub>)alkylcarbonyl; (C<sub>2-6</sub>)alkenyloxycarbonyl; (C<sub>2-6</sub>)alkenylcarbonyl; hydroxy, amino or aminocarbonyl optionally substituted as for corresponding substituents **R<sup>12</sup> as defined** in R<sup>3</sup>;

(C<sub>1-6</sub>)alkylsulphonyl; (C<sub>2-6</sub>)alkenylsulphonyl; or (C<sub>1-6</sub>)aminosulphonyl wherein the amino group is optionally substituted by (C<sub>1-6</sub>)alkyl or (C<sub>1-6</sub>)alkenyl;

or R<sup>6</sup> and R<sup>8</sup> together represent a bond and R<sup>7</sup> and R<sup>9</sup> are as above defined;

or R<sup>6</sup> and R<sup>7</sup> or R<sup>8</sup> and R<sup>9</sup> together represent oxo;

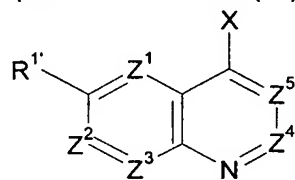
**At page 6, lines 3-9:**

Preferred examples of R<sup>3</sup> include hydrogen; optionally substituted aminocarbonyl; optionally substituted (C~~1-6~~1-4)alkyl; carboxy(C<sub>1-4</sub>)alkyl; optionally substituted aminocarbonyl(C<sub>1-4</sub>)alkyl; cyano(C<sub>1-4</sub>)alkyl; optionally substituted 2-oxo-oxazolidinyl and optionally substituted 2-oxo-oxazolidinyl(C<sub>1-4</sub>alkyl). More preferred R<sup>3</sup> groups are hydrogen; CONH<sub>2</sub>; 1-hydroxyalkyl e.g. CH<sub>2</sub>OH, CH(OH)CH<sub>2</sub>CN; CH<sub>2</sub>CO<sub>2</sub>H; CH<sub>2</sub>CONH<sub>2</sub>; 1,2-dihydroxyalkyl e.g. CH(OH)CH<sub>2</sub>OH; CH<sub>2</sub>CN; 2-oxo-oxazolidin-5-yl and 2-oxo-oxazolidin-5-yl(C<sub>1-4</sub>alkyl).

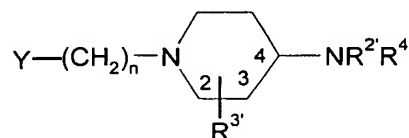
**At page 8, lines 32-36 to page 9, lines 1-31:**

In a further aspect of the invention there is provided a process for preparing compounds of formula (I), and pharmaceutically acceptable derivatives thereof, which process comprises:

reacting a compound of formula (IV) with a compound of formula (V):



(IV)



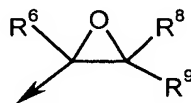
(V)

wherein  $Z^1$ ,  $Z^2$ ,  $Z^3$ ,  $Z^4$ ,  $Z^5$  and  $n$  are as defined in formula (I);  $R^{1'}$ ,  $R^{2'}$ ,  $R^{3'}$  and  $R^{4'}$  are  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  as defined in formula (I) or groups convertible thereto;

and  $X$  and  $Y$  may be the following combinations:

- (i)  $X$  is  $A'-COW$ ,  $Y$  is  $H$  and  $n$  is 0;
- (ii)  $X$  is  $CR^6=CR^8R^9$ ,  $Y$  is  $H$  and  $n$  is 0;
- (iii)  $X$  is oxirane,  $Y$  is  $H$  and  $n$  is 0;
- (iv)  $X$  is  $N=C=O$  and  $Y$  is  $H$ ;
- (v)  $X$  is  $NH_2$  and  $Y$  is  $CO_2W$ ;
- (vi) one of  $X$  and  $Y$  is  $CO_2R^Y$  and the other is  $CH_2CO_2R^X$ ;
- (vii)  $X$  is  $CHR^6R^7$  and  $Y$  is  $CR^8O$ ;
- (viii)  $X$  is  $CR^6=PR^Z_3$  and  $Y$  is  $CR^8O$ ;
- (ix)  $X$  is  $CR^6O$  and  $Y$  is  $CR^8=PR^Z_3$ ;
- (x) one of  $X$  and  $Y$  is  $COW$  and the other is  $NHR^{11'}$  or  $NCO$ ;
- (xi)  $X$  is  $CR^6O$  and  $Y$  is  $NHR^{11'}$  or  $X$  is  $NHR^{11'}$  and  $Y$  is  $CR^8O$ ;
- (xii)  $X$  is  $NHR^{11'}$  and  $Y$  is  $CR^8R^9W$ ;
- (xiii)  $X$  is  $CR^6R^7W$  and  $Y$  is  $NR^{11'}$  or  $O$ ; or
- (xiv)  $X$  is  $CR^6R^7SO_2W$  and  $Y$  is  $H$  and  $n=0$ ;
- (xv)  $X$  is  $NR^{11'}$  and  $Y$  is  $SO_2W$ ;

in which  $W$  is a leaving group, e.g. halogen;  $R^X$  and  $R^Y$  are  $(C_{1-6})$ alkyl;  $R^Z$  is aryl or  $(C_{1-6})$ alkyl;  $A'$  and  $NR^{11'}$  are  $A$  and  $NR^{11}$  as defined in formula (I), or groups convertible thereto; and oxirane is:



wherein  $R^6$ ,  $R^8$  and  $R^9$  are as defined in formula (I);

and thereafter optionally or as necessary converting  $A'$ ,  $R^{1'}$ ,  $R^{2'}$ ,  $R^{3'}$ ,  $R^{4'}$  and  $NR^{11'}$  to  $A$ ,  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$  and  $NR^{11}$ ; converting  $A-B$  to other  $A-B$ , interconverting  $R^1$ ,  $R^2$ ,  $R^3$  and/or  $R^4$ , and/or forming a pharmaceutically acceptable derivative thereof.